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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)		Application Number	09/785,548
		Filing Date	2/20/2001
		First Named Inventor	Koutnikova, Hana
		Group Art Unit	1647
		Examiner Name	Hayes, Robert Clinton
		Attorney Docket Number	ST00005
Sheet	2	of	2

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

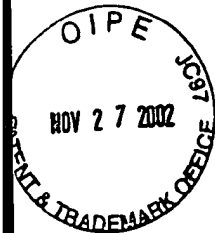
Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published	T ²
not machine	no comp	SPRAUSBERG, P.; Database: Embi Sequence on line; Accession No. AF701000; "EST, H. series cDNA clone IMAGE-2940603 similar to TR. Q14184 DGC2 beta", 6/4/1999.	
PKH		KITADA, T., et al.; Mutations in the parkin gene cause autosomal recessive juvenile parkinsonism; Nature, Vol. 329, April 9, 1998, 605-608.	
		ABBAS, N., et al.; A wide variety of mutations in the parkin gene are responsible for autosomal recessive parkinsonism in Europe; Hum. Mol. Genet., Vol. 8, 1999, 567-574.	
		HATTORI, N., et al.; Molecular genetic analysis of a novel Parkin gene in Japanese families with autosomal recessive juvenile parkinsonism; Ann Neurol., Vol. 6, 1998, 935-941.	
		LUCKING, C., et al.; Homozygous deletions in parkin gene in European and North African families with autosomal recessive juvenile parkinsonism; Lancet, Vol. 352, 1998, 1355-1356.	
		MORETT, E., et al.; A novel transactivation domain in parkin; Trends Biochem Sci., Vol. 24, 1999, 229-231.	
		POLYMERPOULOS, M.H., et al.; Mutation in the alpha-synuclein gene identified in families with Parkinson's disease; Science, Vol. 276, 1997, 2045-2047.	
		SHIMURA, H., et al.; Immunohistochemical and subcellular localization of Parkin protein: absence of protein in autosomal recessive juvenile parkinsonism patients; Ann Neurol., Vol. 45, 1999, 668-672.	
		SUNADA, Y., et al.; Differential expression of the parkin gene in the human brain and peripheral leukocytes; Neurosci Lett., Vol. 254, 1998, 180-182.	
		MELLICK, G.D. et al.; The parkin gene S/N167 polymorphism in Australian Parkinson's disease patients and controls; Parkinsonism and Related Disorders 7, 2001, 89-91.	

Examiner Signature	<i>RP Hayes</i>	Date Considered	2/19/03
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Applicant's unique citation designation number (optional). ² Applicant is to place a check mark here if English language Translation is attached.

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